## LISTING OF CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application.

## What is claimed is:

- 1. (Original) A pharmaceutical composition comprising (i) a first specific binding agent selected from an antibody or a large binding fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin.
- 2. (Previously presented) The composition of claim 1 wherein the first specific binding agent comprises a large binding fragment of an antibody.
- 3. (Previously presented) The composition of claim 2 wherein the large binding fragment of an antibody is an  $F(ab')_2$  or  $F(ab)_2$  fragment.
- 4. (Previously presented) The composition of claim 1 wherein the first specific binding agent is an antibody which is IgG or IgT.
- 5. (Previously presented) The composition of claim 4 wherein the antibody is humanised.
- 6. (Previously presented) The composition of claim 1 wherein the second specific binding agent comprises an Fab, Fab', a single chain (sc) antibody or an FV, VH or VK fragment.

- 7. (Previously presented) The composition of claim 6 wherein the second specific binding agent comprises an Fab or Fab' fragment.
- 8. (Previously presented) The composition of claim 1 wherein the first and/or second binding agents are derived from polyclonal antibodies.
- 9. (Previously presented) The composition of claim 1 wherein the first and/or second binding agents are derived from monoclonal antibodies.
- 10. (Previously presented) The composition of claim 1 wherein at least one of the first or second specific binding agents includes a section corresponding to part of the Fc region of an antibody.
- 11. (Previously presented) The composition of claim 1 wherein the toxin is a Botulinum toxin.
- 12. (Previously presented) The composition of claim 11 wherein the first and second specific binding agents bind at least one of type A, B, C, D, E, F or G botulinum toxin.
- 13. (Previously presented) The composition of claim 12 wherein the composition comprises sets of first and second specific binding agents each set of specific binding agents binding a different one of botulinum toxins A, B, C, D, E, F or G.
- 14. (Previously presented) The composition of claim 1 wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 90:10 to 10:90.

- 15. (Previously presented) The composition of claim 14 wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 70:30 to 30:70.
- 16. (Previously presented) The composition of claim 15 wherein the w/w ratio of the first specific binding agent to the second specific binding agent is in the range of from 60:40 to 40:60.
- 17. (Previously presented) The composition of claim 1 which further comprises a pharmaceutically acceptable carrier or excipient.
- 18. (Previously presented) The composition of claim 1 which is suitable for oral, parenteral, or intranasal administration, or for administration by inhalation or insufflation.
- 19. (Withdrawn) A method for treating the adverse effects of a toxin on a mammal comprising administering to a mammal in thereof a composition comprising (i) a first specific binding agent selected from an antibody or a large binding fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin.

## 20. (Cancelled)

21. (Withdrawn) A method of preventing the effects of a toxin on a mammal, said method comprising administering to a mammal in need thereof, a composition comprising (i) a first specific binding agent selected from an antibody or a large binding

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fragment of an antibody which specifically binds a target toxin, and (ii) a second specific binding agent which comprises a small binding fragment of an antibody which binds said toxin.

22. (Cancelled)